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ARTICLE

Liposome-induced exfoliation of graphite to few-layer graphene dispersion with antibacterial activity

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Few-layer graphene aqueous dispersions are obtained by exploiting liposomes as effective exfoliating agent for graphite. Raman measurements evidence the presence in the samples of non-oxidized double layer graphene as well as amphiphilic phospholipid molecules organized in bilayers. TEM analyses confirmed that the obtained homogeneous graphene nanosheets are embedded in the liposomal bilayer. The as-prepared graphene aqueous dispersion is stable for days and demonstrates to behave as a significant antibacterial agent towards both Gram-positive (*Staphylococcus aureus*) and Gram-negative (*Escherichia coli*) strains, with a reduction in the growth of *S. aureus* and *E. coli* as high as 60 and 78%, respectively.

Introduction

In recent years a growing interest within the scientific community has been devoted to the study of carbon allotropes and their applications. Among them, graphene and its derivatives, due to their unique physicochemical properties, have aroused interest¹⁻³ in many research fields. As a matter of fact, applications in electronic⁴⁻⁷ and photonic devices,⁸⁻¹⁰ clean energy,^{1,11,12} energy storage¹³ and sensors^{14,15} have been well demonstrated. In addition, graphene-based materials appear as promising scaffolds in biomedicine.^{1-3,16-18}

For application in biomedicine one of the most important and fundamental goal to be achieved is to make graphene soluble in water. Up to now, most of the research on graphene in the biomedical field has been focused on the production and characterization of hydrophilic graphene oxide (GO), i.e. graphene functionalized with hydroxyl, epoxy and carboxyl groups that render graphene suitable to interface with biological systems. As a matter of fact, the functionalization, renders GO-based material chemically versatile templates of high surface-to-volume ratio and favours the realization of GO-based drug delivery vehicles,¹⁹⁻²¹ biosensors,²²⁻²⁶ imaging agents^{19,27} and electromechanical devices for monitoring cellular responses.²⁸

Besides GO discloses good biocompatibility against animal cells,²⁹ while stimulating human mesenchymal stem cells to differentiate into osteoblasts,^{30,31,32} adipocytes³⁰ and

myoblasts,^{33,34} inducing the differentiation of preosteoblasts into osteoblasts,^{35,36} of neuronal stem cells into neurons³⁷ and pluripotent stem cells into endodermal lineage.³⁸ For this reason GO-base materials have been studied for tissue and osteoregeneration.^{39,40,41} GO demonstrated to exhibit strong antibacterial activity towards both Gram-positive and Gram-negative bacteria.^{42,43}

The functionalization of graphene to get GO has revealed undisputed advantages, but it involves the breakdown of the continuous honeycomb backbone of pristine (non-functionalized) graphene compromising several of the peculiar properties of the original material.

In order to preserve graphene structural integrity and to use an absolutely green process of exfoliation and functionalization, inspired by Samori *et al*⁴⁴ that demonstrates the excellent ability of fatty acids to exfoliate graphite in organic solvents and Titov's molecular dynamics simulation⁴⁵ that showed the theoretical insertion of a "graphene sheet in the hydrophobic interior of biological membranes", in the present study we succeed to obtain nanosheets of graphene sandwiched between phospholipid alkyl chains by simply sonicating graphite into liposomal suspensions. The as-prepared aqueous dispersions demonstrate to solubilize as much as 161 $\mu\text{g mL}^{-1}$ few-layer graphene in line with dispersions obtained with the use of classical surfactants such as sodium cholate but three times more concentrated than dispersions obtained by sodium benzenedodecyl sulfonate.⁴⁶ The advantage of the present dispersions is that flakes are relatively small i.e. (in the nanometric range) and ready to be used for different biomedical applications.

Considering that (i) resistance to antibiotics represents the major cause of treatment failure in bacterial infections, (ii) alternative proposals are needed due to overuse or misuse of common antimicrobials that compromises the therapeutic effect of traditional treatments⁴⁷ and (iii) the proved antibacterial activity of the covalently functionalized oxidized

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analogue GO, we propose the as-prepared liposomal graphene-loaded aqueous dispersion as innovative efficacious antibacterial system to overcome the alarming bacterial resistance increase.^{48,49} Therefore, in the present study, the antibacterial activity of the as-prepared non-covalently functionalized graphene aqueous dispersion against a Gram-positive (*Staphylococcus aureus*) and a Gram-negative (*Escherichia coli*) strain is evaluated and compared with that of the analogous GO.

Results and discussion

Graphene (G) dispersion inside liposome bilayer in aqueous solution

Homogeneous aqueous suspensions of graphene-loaded liposomes (LIPO-G) were prepared through sonication of graphite in the presence of liposomes. The exfoliation was monitored via UV-vis absorption spectra in the 200-800 nm wavelength range. The observed flat and featureless spectrum is typical of the exfoliated graphitic material,⁵⁰ absorption from liposomes being negligible above 400 nm (see Figure 1).

Such observation allowed to use the suspension absorbance at 660 nm as a measure of the concentration of exfoliated graphene loaded in the liposome bilayer. Graphene quantity was estimated from the absorbance at 660 nm by using the extinction coefficient ($\epsilon = 1390 \text{ mg}^{-1} \text{ mL m}^{-1}$) previously determined in water.⁵⁰ With the designed protocol of graphene exfoliation and dispersion inside the liposomal bilayer, a yield of $15.6 \pm 1.0 \%$ in PBS ($[G] = 0.161 \pm 0.010 \text{ mg mL}^{-1}$) and $13.2 \pm 0.4 \%$ in milli-Q water ($[G] = 0.124 \pm 0.010 \text{ mg mL}^{-1}$) was obtained. Instead a graphene entrapment efficiency of the investigated liposomes (i.e. $[\text{weight of G}]/[\text{weight of phospholipids}] \%$) of $1.9 \pm 0.1 \%$ could be calculated in PBS.

Dynamic light scattering experiments revealed that the as-prepared LIPO-G had diameters in the 240-310 nm range (i.e.

$238 \pm 6 \text{ nm}$ for samples in PBS and $307 \pm 3 \text{ nm}$ for samples in milli-Q water), which is the typical feature of large unilamellar vesicles (LUV). A diameter of $222 \pm 7 \text{ nm}$ was measured for POPC liposomes without graphene.

ζ -Potential measurements showed that LIPO-G had an average negative ζ -potential of $-15.2 \pm 0.3 \text{ mV}$. Pure POPC liposomes without graphene had a comparable ζ -potential ($-15.4 \pm 0.8 \text{ mV}$). Because pristine graphene ζ -potential in water is $-21.8 \pm 0.05 \text{ mV}$,^{51,52} these evidences pointed to perfectly sandwiched graphene in the liposomal bilayer.

Stability of graphene-loaded liposome

Liposome dimensions (and relevant polydispersity index)⁵³ were checked over 48 h (see Figure S1 in the ESI) and the corresponding dispersions demonstrated to be relatively stable over time. It is worth noting that polydispersity for the obtained LIPO-G ranged between 0.25-0.30, underlying a fairly homogeneous liposomal population once considered that sonication was the only homogenization step performed.⁵³

The ζ -potential is often considered a further measure of the colloidal stability of a suspension, in particular ζ -potential higher than +15 or lower than -15 mV are indicative of stable colloidal samples.⁵⁴ The ζ -potential for the obtained samples ranged between -15.7 and -14.8 mV during 48 h (see Figure S2 in the ESI).

In view of a potential administration in a biological environment another parameter to consider for materials intended to be used in biological samples is the stability upon dilution. As reported in Figure S3 in the ESI, the stability of the diluted samples overwhelmed that of the as-prepared samples either in terms of dimensions and polydispersity index.

Evidence of graphite exfoliation

Raman spectroscopy was used to confirm the exfoliation of graphite in few-layer graphene by exploiting POPC phospholipids. The Raman spectra of graphene showed the G peak located at $\sim 1580 \text{ cm}^{-1}$ and the 2D peak at $\sim 2700 \text{ cm}^{-1}$ (see Figure 2). The small D peak, located at $\sim 1350 \text{ cm}^{-1}$ is due to first-order zone boundary phonons and is indicative of the presence of some defects in the exfoliated graphene (see Figure 2). It is accepted that Raman could be used to quantify the degree or type of defects of the exfoliated graphene and to determine the number of layers by monitoring the shape, width, and position of the 2D peak.⁵⁵

The Lorentzian fitting of the 2D Raman band allowed to ascertain that treatment with liposomal suspension caused a good exfoliation of graphite into a few-layer graphene.⁵⁶ As a matter of fact, fitting of 2D peaks recorded over 14 different spots highlighted the presence of double-layer graphene (see Figure 3 as an example and Figure S4 and S5 in the ESI).

Raman spectroscopy is also particularly useful for investigating fluidity and conformational order in liposome bilayer⁵⁷⁻⁶⁰ as, in POPC liposomes, the assignment of the peaks is well-established.⁶¹⁻⁶³ According to the literature^{63,64} the I_{2895}/I_{2855} , the I_{2931}/I_{2895} and I_{1125}/I_{1086} Raman intensity ratios can be

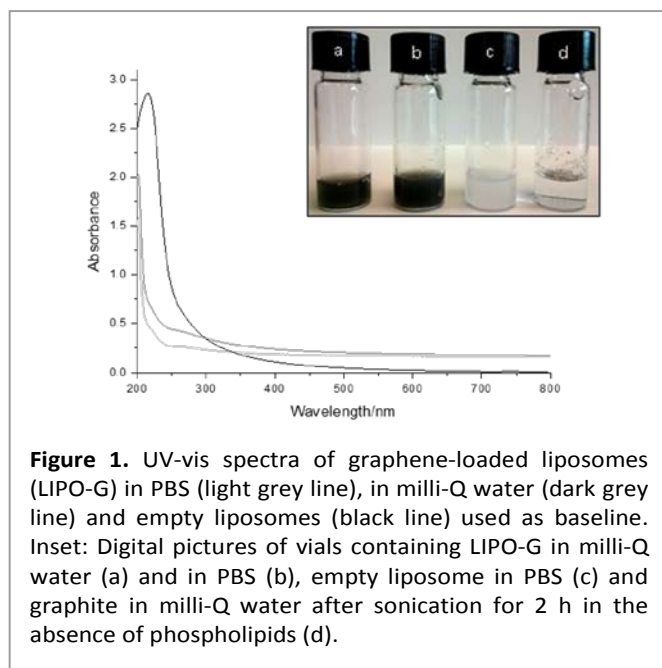


Figure 1. UV-vis spectra of graphene-loaded liposomes (LIPO-G) in PBS (light grey line), in milli-Q water (dark grey line) and empty liposomes (black line) used as baseline. Inset: Digital pictures of vials containing LIPO-G in milli-Q water (a) and in PBS (b), empty liposome in PBS (c) and graphite in milli-Q water after sonication for 2 h in the absence of phospholipids (d).

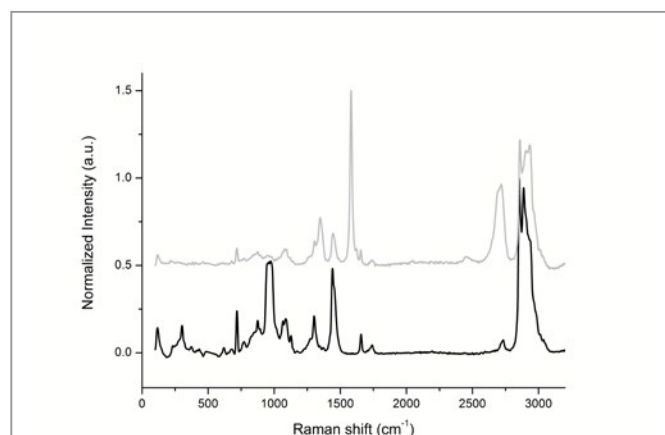


Figure 2. Raman spectra of liposomal suspension (black line) and LIPO-G (grey line). The peak at 1000 cm^{-1} is due to the silicon wafer.

considered good indicators of the molecular order of the lipid bilayer. Indeed, the I_{1125}/I_{1086} Raman intensity ratio is related to the average number of “trans” bonds in the acyl chain, thus giving a measure of the order in the intra-chain structure. On the contrary, the I_{2895}/I_{2855} Raman intensity ratio, depending on the vibrational coupling between the adjacent POPC alkyl chains, reveals the lateral interactions between lipid chains whereas I_{2931}/I_{2895} reflects both the inter- and intra-chain order/disorder features.

The inclusion of graphene into the bilayer showed a slight increase of the order of the liposomal system as demonstrated by the increase of both I_{2931}/I_{2895} (from 0.87 to 0.89) and I_{2895}/I_{2855} (from 0.84 to 1.05) values. The latter intensity ratio increase was unexpected because interactions with guests penetrating the bilayer is generally associated with a I_{2895}/I_{2855} decrease.⁶⁵ Likely the presence of graphene did not hamper inter-chains interactions and therefore bilayer self-assembly (i.e. the I_{2931}/I_{2895} ratio did not vary), but favoured additional interactions between alkyl chains of POPC molecules and graphene layers.^{66,67} Such evidences pointed to a prevailing

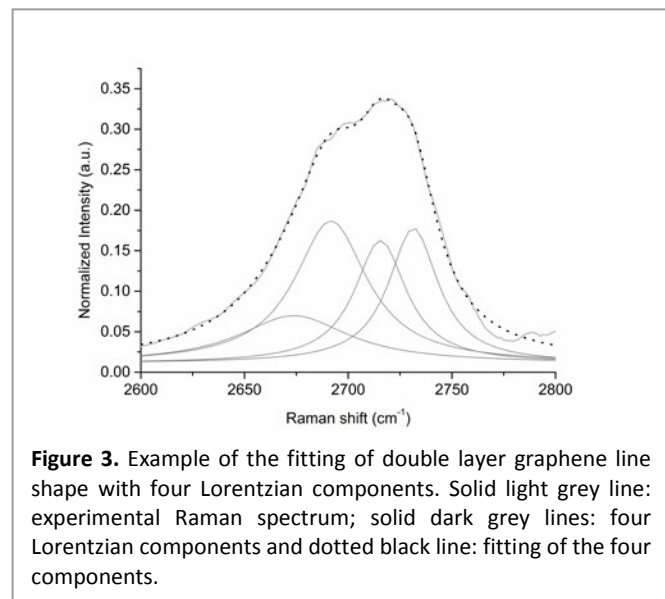


Figure 3. Example of the fitting of double layer graphene line shape with four Lorentzian components. Solid light grey line: experimental Raman spectrum; solid dark grey lines: four Lorentzian components and dotted black line: fitting of the four components.

arrangement of exfoliated graphene inside the core of the bilayer in perfect agreement with coarse-grained dynamic simulations that showed graphene perfectly sandwiched into POPC bilayers and almost not affecting its thickness.⁴⁵ Graphene bilayers slowly diffused in the membrane interior, but the composite system stayed stable over time. Accordingly, the I_{1125}/I_{1086} Raman intensity ratio decreased thus denoting a decrease in the average number of “trans” bonds in the alkyl chains of POPC as a consequence of folding of the alkyl chains to promote van der Waals interactions between alkyl chains and graphene surface.

In order to further confirm the exfoliation degree of graphene, we performed AFM analysis. Figure 4 reports AFM images of LIPO-G and the corresponding samples cross section along the cross line. The samples, prepared by drop-casting a dilute LIPO-G solution onto a silicon wafer, were then dried under vacuum and washed with methanol to get rid of excess liposomes and phospholipids. The cross section AFM view showed that the minimum height of the liposome coated sample is ca. 3-4 nm. By considering that (i) the sample is formed of graphene covered with variable amounts of phospholipids and is therefore a difficult sample to be analyzed by AFM due to the dependence of the AFM measurements on the roughness and cleanness of the surface,⁶⁸ (ii) the thickness of a phospholipid bilayer is around 0.4 nm,⁶⁹ (iii) stabilizers cannot touch the surface of graphene directly^{68,70} and a gap exists between phospholipids and graphene, and (iv) AFM measurements depends strongly on the substrate used for graphene deposition and are distorted by the variability of substrates,^{68,71,72} these data supported the formation of a few-layer graphene, likely not more than 2-3 layers.

Figure 5 reports typical TEM images of the analyzed sample. The results showed that vesicles belong to the class of LUVs with a mean diameter of about 150-180 nm for empty liposomes (Figure 5A) and 240-280 nm for LIPO-G (Figure 5B-5D). While LIPO-G LUVs were easily observed without the use of vesicle staining, empty LUVs were difficult to focalize and visualize on the TEM grid. The loading of graphene did not significantly alter

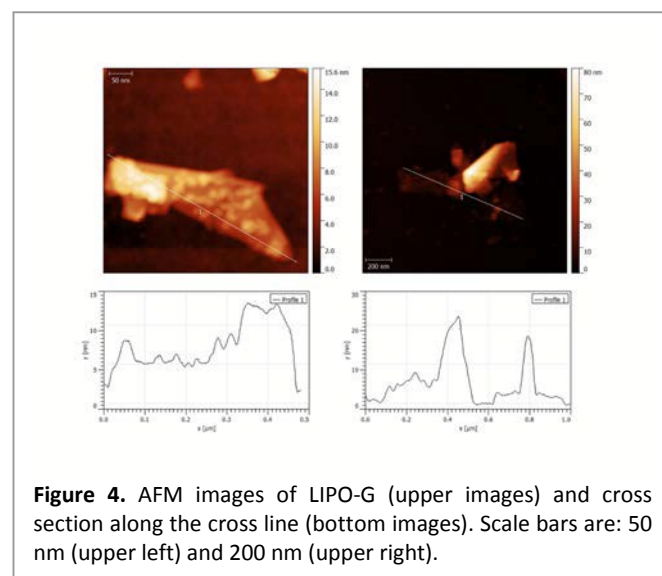


Figure 4. AFM images of LIPO-G (upper images) and cross section along the cross line (bottom images). Scale bars are: 50 nm (upper left) and 200 nm (upper right).

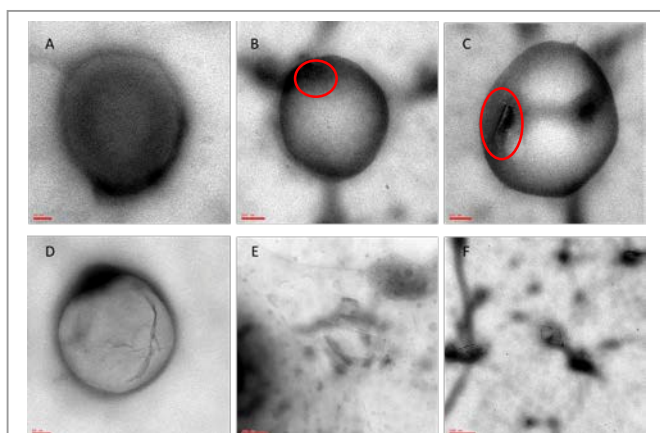


Figure 5. TEM images of empty liposome magnified at 140000 \times (A), LIPO-G magnified at 85000 \times (B and C) and 140000 \times (D) and graphene sheets magnified at 50000 \times (E and F). Scale bars are: 200 nm for 50000 \times images, 100 nm for 85000 \times images and 50 nm for 140000 \times images. All samples were performed in milli-Q water. Red circles indicate partially hindered graphene sheets.

LUV morphology and, during the experiments, LIPO-G LUVs appeared to be able to stand electron beam for longer time with respect to empty liposomes. Figure 5B-5D clearly evidenced the presence of graphene sheets inside the liposomes although, due to the need to eliminate the solvent, the real disposition of graphene with respect to the not dried bilayer was difficult to envisage. Only very few graphene flakes were observed outside the liposomes deposited onto the TEM grid (Figure 5E and 5F). In order to ascertain that graphene was inserted inside the liposomes we destroyed the liposomes by addition of 10 μ L of a concentrated Triton X-100 aqueous solution to the liposomal solution. The precipitation of graphene flakes was detectable by eye (see Figure S6 in the ESI). The so obtained solution was deposited onto a copper grid and visualized by TEM. Several very small (ca. 100 nm) graphene flakes could be detected this time on the grid (Figure 6).

The size and size distribution of LUVs in the as-prepared dispersion were checked also by DLS and the results were in good agreement with TEM measurements (see previously reported data).

Antimicrobial activity

Staphylococcus aureus and *Escherichia coli* were used to evaluate the antibacterial effect of graphene oxide (GO) and LIPO-G at the same concentration, 50 μ g mL⁻¹. The saline (SF) and liposome suspension (LIPO) without graphene were used as the controls. The antibacterial activity of LIPO-G, compared to that recently reported⁷³ for GO with respect to the control LIPO and SF, respectively, after 2 h contact is shown in Figure 7.

The number of colony forming units of *S. aureus* ATCC 29213 and *E. coli* ATCC 8739 was slightly reduced in the presence of both GO and LIPO-G when compared to the CFUs detected with saline and LIPO. This reduction was significant ($P < 0.05$) in *S. aureus* after GO and LIPO-G treatment vs. SF, as well as in *E. coli* after GO and LIPO-G treatment vs. SF (Figure 7A). The reduction

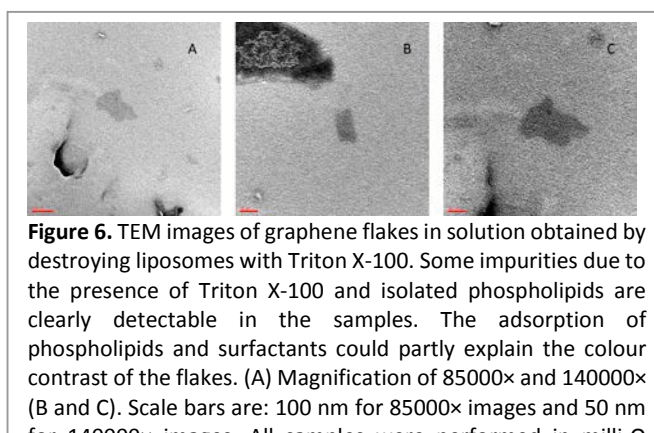


Figure 6. TEM images of graphene flakes in solution obtained by destroying liposomes with Triton X-100. Some impurities due to the presence of Triton X-100 and isolated phospholipids are clearly detectable in the samples. The adsorption of phospholipids and surfactants could partly explain the colour contrast of the flakes. (A) Magnification of 85000 \times and 140000 \times (B and C). Scale bars are: 100 nm for 85000 \times images and 50 nm for 140000 \times images. All samples were performed in milli-Q

of CFUs is clearly shown on TSA and LB agar plates (Figure 7B) where the count of colonies was performed.

Figure 8 displays the reduction of bacterial growth with the drop methodology. After 24 h of incubation in the presence of GO and LIPO-G, both *S. aureus* and *E. coli* colonies were reduced with respect to the controls.

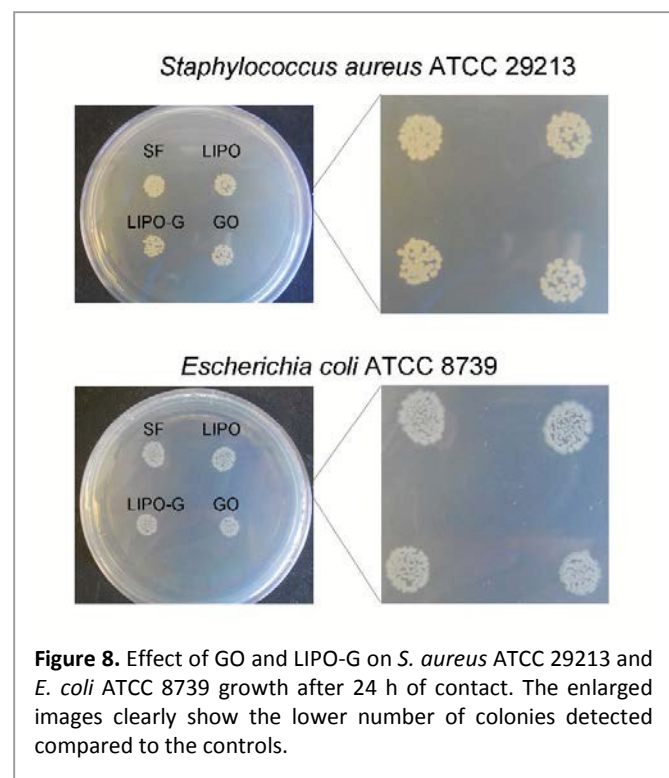
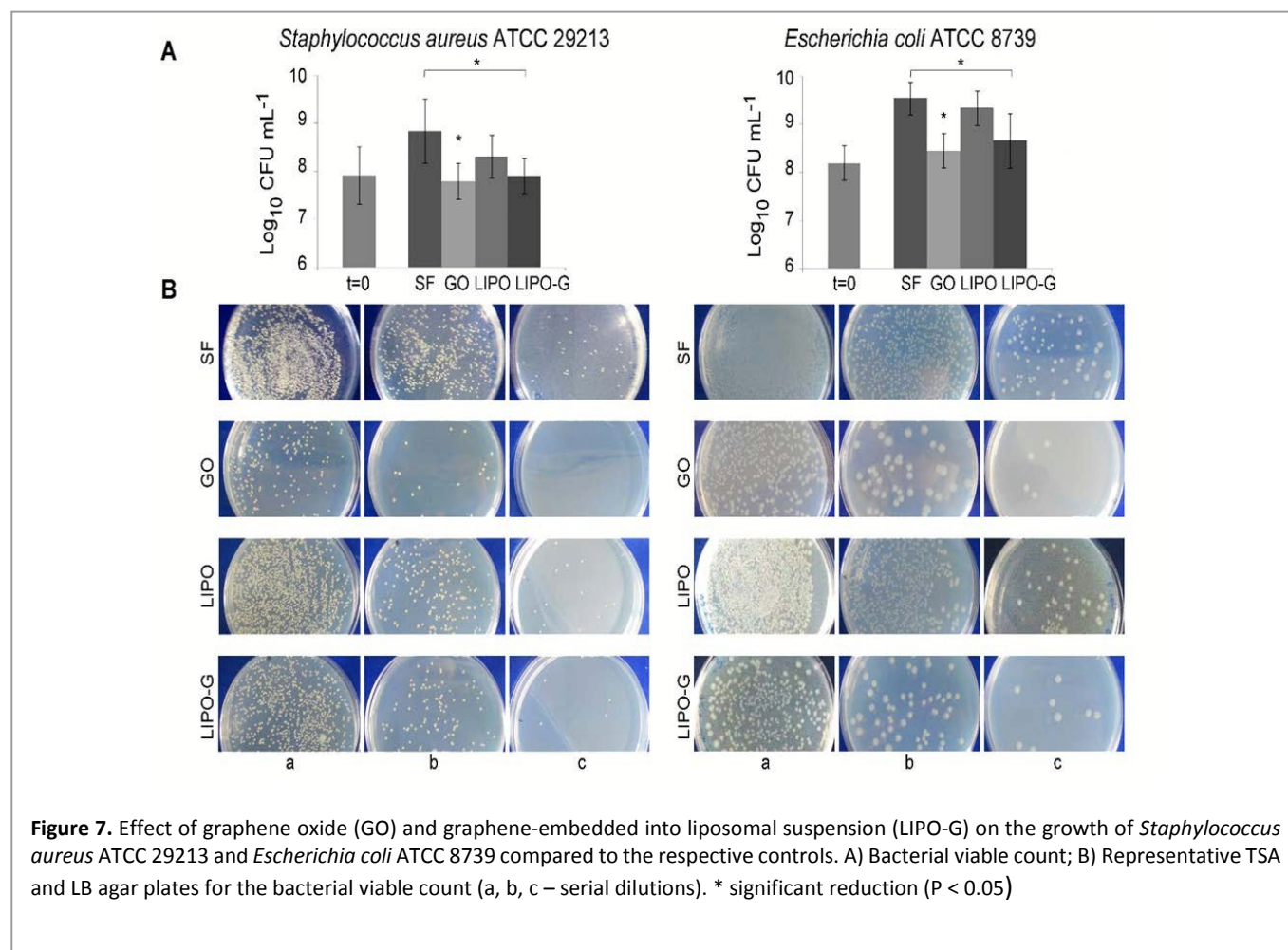
The microscopic analysis of bacterial colonies confirmed the goodness of the reading. In fact, the observed bacteria do not aggregate to each other underlying that each counted colony derived from one singular bacterial cell (see Figure S7 in the ESI). The percentage of reduction of bacterial growth in the presence of GO and LIPO-G, obtained by colonies count (see Figure 7), is reported in Table 1.

Values of bacterial growth reduction of 91.0% for *S. aureus* and 91.9% for *E. coli* were recorded after treatment with GO with respect to the control (SF) in agreement with literature data.⁷³ Instead reduction of 60.1% for *S. aureus* and 78.5% for *E. coli* were recorded after treatment with LIPO-G with respect to the control (LIPO). It is worth noting that the treatment with LIPO-G, with respect to SF, induced a reduction of 88.4% and 86.5% respectively, for *S. aureus* and *E. coli*, highlighting an antimicrobial activity comparable to that reported for the hydrophilic GO. These data are particularly interesting because they point out the ability of liposomes to mediate the antimicrobial activity of the hydrophobic and non-covalently functionalized graphene.

No significant selectivity of LIPO-G is detected for Gram-positive or Gram-negative bacteria although an almost complete inhibition of growth of Gram-positive is monitored with respect to $t = 0$. This datum is particularly interesting as GO evidenced instead a killing effect.

As far as the mechanism of antibacterial activity is concerned, it is likely, following previously published papers,^{42,43} and the observed affinity of pure liposomes and LIPO-G for the external membrane of the cell wall of Gram-negative, that a damage of the membrane was involved. Indeed, pure liposomes demonstrated to stimulate mostly the growth of Gram-negative with respect to Gram-positive bacteria due to the similar phospholipidic composition of the external membrane of their cell wall. Analogously, graphene non-covalently functionalized with phospholipids reduced the growth of Gram-negative twice as much as that of Gram-positive.

The bacterial viability is reported, by using Live/Dead staining, in Figure 9. *Staphylococcus aureus* and *Escherichia coli* cells



were uniformly viable with a green fluorescence and only few red dead cells.

Taken together these data confirmed that the main effect obtained after contact with GO and LIPO-G was a bacterial growth inhibition and a very low killing effect. In fact, after 2 h of incubation, the bacteriostatic effect was clearly detectable by the smaller number of bacteria for field with respect to the bacterial population in the controls. Moreover, these treated bacteria appeared almost totally viable with only few dead red cells strongly suggesting a bacteriostatic effect of GO and LIPO-G on both *S. aureus* and *E. coli*.

Table 1. Percentage of *Staphylococcus aureus* ATCC 29213 (SA) and *Escherichia coli* ATCC 8739 (EC) growth reduction after treatment with GO and LIPO-G when compared with the respective controls (SF and LIPO).

Strain	CFU mL ⁻¹		% R ^a	CFU mL ⁻¹		% R ^b	% R ^c
	t = 0	SF		LIPO	LIPO-G		
SA	8.21×10 ⁷	6.97×10 ⁸	6.30×10 ⁷	91.0	2.02×10 ⁸	8.06×10 ⁷	60.1 88.4
EC	1.55×10 ⁸	3.45×10 ⁹	2.81×10 ⁸	91.9	2.16×10 ⁹	4.65×10 ⁸	78.5 86.5

^a) Bacterial growth reduction GO vs. SF; ^b) Bacterial growth reduction LIPO-G vs. LIPO; ^c) Bacterial growth reduction LIPO-G vs. SF

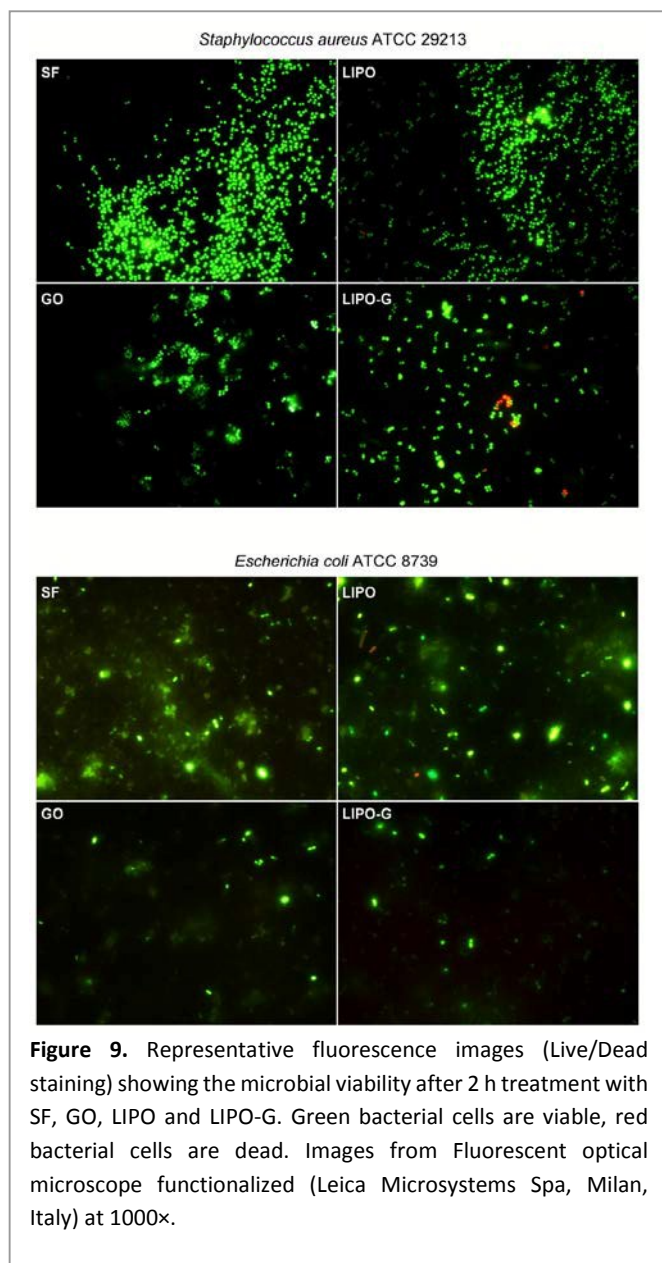


Figure 9. Representative fluorescence images (Live/Dead staining) showing the microbial viability after 2 h treatment with SF, GO, LIPO and LIPO-G. Green bacterial cells are viable, red bacterial cells are dead. Images from Fluorescent optical microscope functionalized (Leica Microsystems Spa, Milan, Italy) at 1000 \times .

Experimental

Materials

Graphite powder (99.99 %, 200 mesh) was purchased from Alfa Aesar, 1-palmitoyl-2-oleyl-*sn*-glycero-3-phosphocoline (POPC) was purchased from Avanti Polar Lipids, Inc. (Alabaster, AL). Graphene oxide used for comparative antibacterial activity determination (GO) was prepared by using the modified Hummers method.^{40,74}

Unless otherwise stated, reagents were of analytical grade and used as received. All aqueous solution were prepared with Ultrapure Milli-Q water (electric resistance > 18.2 M Ω cm⁻¹) from a Millipore Corp. model Direct-Q 3 system.

Apparatus

Sonication was performed by using an ultrasonic bath (Transsonic 310 Elma, 35 kHz). UV-vis spectra were recorded on a Jasco V-550. Measurement of vesicle size and ζ -potential values were performed by using a 90Plus/BI-MAS Zeta Plus multiangle particle size analyzer (Brookhaven Instruments Corp.). Raman spectra were recorded with an Invia Renishaw microspectrometer (50 or 100 \times) and a laser source at 532 nm (power 5 %, 3 or 5 accumulations/measurement). TEM observations were carried out on a Zeiss Electron Microscope M109 operated at 80 kW working voltage. Surfaces were imaged with a scan rate from 0.1 to 1.95Hz. Images were acquired by Gatan camera (U.S.) and processed by Nanoscope Analysis software. AFM analysis was performed on a Nanoscope V (Digital Instruments Metrology Group, model MMAFMLN) in tapping mode in air at room temperature, using a n-type silicon μ mash[®] SPM probe (HQ:NSC15/AL BS) with tip height 12-18 μ m, cone angle < 40 $^\circ$ (Resonant frequency 325 kHz, force constant of \sim 40 N/m). The collected images were then analyzed with WsXm 4.0 software (Nanotec Electronica S. L.) and Gwyddion 2.39.

Preparation and processing of graphene-loaded liposome

Liposomes were prepared according to the following protocol. 40 mg of POPC, dissolved in 4 mL of chloroform, were put in a round-bottom flask and dried in a rotary evaporator under reduced pressure at 40 $^\circ$ C to form a thin lipid film on the inside wall of the flask. The phospholipidic film was kept at 4 $^\circ$ C overnight before rehydration with 5 mL of Milli-Q water or PBS buffer (pH 7.4, 304 mOsm) in order to obtain a 0.8 % w/w aqueous solution of phospholipid, concentration that is above POPC critical aggregation concentration value (0.1 % w/w). 5 mg of graphite powder was added to the resulting liposome suspension and sonicated for 2 h. The resulting mixture was centrifuged at 5000 rpm for 15 min to sediment unexfoliated particles or thick flakes of graphite and the supernatant, which was the final LIPO-G suspension, was collected.

Exfoliation yield and encapsulation efficiency for LIPO-G were calculated via UV-Vis spectrophotometry by determining the amount of graphene present into the supernatant after liposomal centrifugation.

The liposomal suspension was characterized by using UV-vis spectroscopy, dynamic light scattering (DLS), ζ -potential, transmission electron microscopy (TEM), Raman spectroscopy and AFM. The samples for Raman characterization were obtained by dropping the liposomal suspension on a wafer's surface (we used n-type doped prime SiO₂ wafers) and leaving the sample in the oven at 40 $^\circ$ C for 24 h in order to dry the aqueous solution. TEM samples were prepared by dropping a small drop of liposomal suspension (about 5 μ L) on a carbon coated copper grid. Solvent was removed by keeping the grid at 30 $^\circ$ C for 60 h into an incubator in the presence of silica gel in order to ensure a total absence of moisture. The choice of using a low drying temperature for a relatively long time is connected with the need to avoid high temperature and thus graphene sheets folding. AFM samples were prepared by drop-casting the solution onto silicon wafer substrates, drying under vacuum and washing with methanol the sample to remove excess liposomes and phospholipids.

Bacterial culture

Staphylococcus aureus ATCC 29213 and *Escherichia coli* ATCC 8739 were used in the present study. *Staphylococcus aureus* and *E. coli* were cultured in Trypticase Soy Broth (TSB, Liofilchem, Italy) and Luria Bertani broth (LB, Oxoid, Italy) medium, respectively, and incubated at 37 $^\circ$ C overnight in aerobic condition. Cultures were diluted 1:10 (v/v) in TSB and LB medium, respectively and refreshed for 2 h at 37 $^\circ$ C in an orbital shaker in aerobic condition to obtain an

exponential growth phase with a homogeneous bacterial population. Subsequently, the broth-cultures were washed in PBS (pH 7.2) and adjusted to optical density (Ab_{560}) of 0.12 corresponding approximately to 5×10^7 CFU mL^{-1} and used for the experiments.

Antimicrobial activity

The standardized broth-cultures of *S. aureus* ATCC 29213 and *E. coli* ATCC 8739 were incubated with the same final concentration of 50 $\mu g mL^{-1}$ of GO, used as a comparison for the antibacterial activity of pristine graphene, and graphene-embedded into liposomal suspension (LIPO-G); as controls, bacteria were incubated with fresh isotonic saline solution (SF) and an equally concentrated POPC liposomal suspension (LIPO) at 37 °C under 250 rpm in an orbital shaker for 2 h. The loss of bacterial viability was evaluated by counting the colony forming units (CFUs) through the Colony Counter Star-Count STC 1000 (VWR International PBI Srl, Via San Giusto, Milan, ITALY). Briefly, series of 10-fold cell dilutions (100 μL) were spread onto Trypticase Soy Agar (TSA, Liofilchem, Italy) plates for *S. aureus* and onto LB agar (LB, Oxoid, Italy) plates for *E. coli* and incubated for 24 h at 37 °C. The cell growth inhibition was detected comparing the colony counts between GO vs. SF and LIPO-G vs. LIPO. All experiments were performed in duplicate, and repeated for at least three independent experiments. The significance of the differences recorded in the experiments performed with SF, GO, LIPO, LIPO-G were evaluated using Student's t-test. Probability levels < 0.05 were considered statistically significant.

The inhibition of CFUs formation was also performed by dropping 10 μL of tested substances (i.e. SF, GO, LIPO, LIPO-G) on agar plate with 10 μL drop of standardized bacterial suspensions. Plates were incubated at 37 °C for 24 h (see Table 1).

After SF, GO, LIPO, LIPO-G treatment for 2 h, the bacterial viability was also evaluated with Live/Dead Kit (Molecular Probes Inc., Invitrogen, Italy) as indicated by the manufacturer and visualized under a fluorescent Leica 4000 DM microscope (Leica microsystems Spa, Milan, Italy). Ten fields of view randomly chosen for each slide were observed. Microscopic observations were repeated for three independent experiments.

Conclusions

We present a facile and prompt exfoliation protocol for graphite by exploiting 2 h sonication of graphite in POPC large unilamellar vesicles aqueous solution. Spectrophotometry was used to quantify the exfoliated graphene. Raman spectroscopy demonstrated the prevailing formation of double-layer graphene and AFM confirmed the formation of few-layer graphene. TEM pointed out that sonication allows the formation of nanometric few-layer graphene sheets embedded into liposomes. The corresponding aqueous solution demonstrated to form solution stable for hours and able to perform a significant antibacterial activity. In particular, data evidenced that liposome-embedded graphene reduces the growth capability of bacteria. In particular, it almost completely inhibits microbial growth of Gram-positive bacteria whereas Gram-negative bacteria growth is reduced to a mere three fold growth.

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Textual for graphical abstract

Few-layer graphene was obtained by exploiting liposome induced exfoliation of graphite. The obtained graphene nanosheets are homogenous in terms of dimensions and the non-covalent functionalization with phospholipids confers graphene a hydrophilic character. The obtained aqueous dispersions are stable for 48 h and demonstrate antimicrobial activity.

